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(54) Title: TITANIA CATALYSTS, THEIR PREPARATION AND USE IN FISCHER-TROPSCH SYNTHESIS			
(57) Abstract <p>Supports for Fischer-Tropsch catalysts with increased strength and attrition resistance are formed by incorporating both <u>silica</u> and <u>alumina into a support comprised primarily of titania</u>; whereupon Fischer-Tropsch active metals can be composited with the support; the catalysts being particularly useful in slurry reactions.</p>			

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TITANIA CATALYSTS, THEIR PREPARATION AND USE IN FISCHER-TROPSCH SYNTHESIS

FIELD OF THE INVENTION

This invention relates to titania containing supports, their preparation, and use as catalysts (with added metal(s)) for Fischer-Tropsch hydrocarbon synthesis. More particularly, this invention relates to material that can withstand the high water partial pressures often encountered in hydrocarbon synthesis processes, for example, slurry based processes.

BACKGROUND OF THE INVENTION

United States Patent No. 5,140,050 describes improvements in the preparation of supports useful in the preparation of Fischer-Tropsch catalysts, and the unexpected results obtained from the use of these catalysts in the Fischer-Tropsch process. As useful as these improved catalysts and supports have been, the operating conditions for Fischer-Tropsch synthesis, particularly the relatively high water partial pressures that occur as a result of the Fischer-Tropsch reactions, have led to the weakening of the catalysts and the formation of excessive fines in the reaction mixture. The fines have several deleterious effects including the clogging of lines, reduced catalyst efficiency, and loss of catalyst through filters, as well as clogging of filters. Consequently, there is a need for the development of a catalyst that can retain its integrity -- and thereby its efficiency -- under steaming conditions such as are present during the Fischer-Tropsch process. In particular, the severity of slurry operations combined with the steaming conditions requires a catalyst of high attrition resistance.

SUMMARY OF THE INVENTION

In accordance with this invention, both silica and alumina are employed as binders for a titania containing support to achieve the integrity required of a catalyst used in Fischer-Tropsch synthesis. The catalyst is formed by dispersing one or more metals active for Fischer-Tropsch synthesis, e.g., Group VIII metals such as cobalt or ruthenium, over the surface of the support. Thus, a high-strength -- as measured by attrition resistance -- catalyst is formed that maintains its integrity under conditions of relatively high water partial pressure at elevated temperatures, e.g., 175-400°C, used in Fischer-Tropsch processes.

U.S. Patents #5,140,050 and 4,992,406 disclose porosity improvements for titania containing supports by virtue of the use of alumina or zirconia or silica binder materials -- in that order of preference. The invention described herein, however, discloses that the use of silica, a less preferred material than alumina as a binder, together with alumina, produces a titania containing material with greater attrition resistance, either in a dry state or under steaming conditions, than either alumina or silica alone. Consequently, there is a seeming anomaly in the fact that the addition of a less preferred material to a composition containing titania and alumina results in an even stronger support material.

The mechanism by which the synergy of alumina and silica produce high strength titania containing material is unclear. However, while not wishing to be bound by any particular mechanism, we theorize that in the finished, calcined support small silica particles occupy positions at the interface between larger alumina particles and much larger titania particles, and serve as a

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kind of chemical bonding agent. Thus, silica crystallites can partially merge with both alumina and titania crystallites to form an essentially continuous crystalline phase. The silica acts as a binding agent between the alumina and the titania, and the steaming either during calcination or Fischer-Tropsch synthesis, may further the formation of these mixed boundary phases of silica with alumina and silica with titania. Further, alumina by itself does not mix well with titania and only physical forces are relied upon to bind the particles. Steaming may well disrupt these physical forces, possibly by sintering the alumina and forming larger alumina particles that are less able to provide binding qualities. Silica by itself is not a good binder because of its affinity for titania which causes the silica to lose much of its separateness.

DESCRIPTION OF THE DRAWINGS

Figure 1 is a plot of the sonic attrition of various samples where the ordinate represents wt% fines of <25 microns and the abscissa is minutes in an ultrasonic bath. Curves A, B, C and D represent titania supports with steamed alumina, fresh alumina, steamed silica-alumina, and fresh silica-alumina binders, respectively.

DESCRIPTION OF PREFERRED EMBODIMENTS

The titania containing support is a particulate material preferably containing at least about 50 wt% titania, more preferably at least about 80 wt% titania and preferably has a rutile:anatase ratio of at least about 1.9. This material is admixed with suitable amounts of silica and alumina acting as binder materials, where the binder makes up less than about 30 wt%, preferably less than about 20 wt%, more preferably about 3-20 wt%, still more preferably 4-15 wt%, yet more preferred 5-10 wt% of the

total support. The silica and alumina binder mixture may contain 50 wt% or less silica, preferably about 3-50 wt% silica, more preferably 5-35 wt% silica.

The support is typically formed into porous particles with essentially spherical or cylindrical shapes by the well-known methods of extrusion, pilling, tableting, spray-drying, etc. A preferred method is spray-drying, in which a suitable aqueous slurry of titania and binder materials is atomized into a chamber purged with heated air. Spray-drying produces a spherical support with a size range of about 20 to 120 microns, which is well suited for use in slurry Fischer-Tropsch processes.

In order to achieve the benefits of porosity and strength, binder components are mixed with the titania starting material before the forming operation. They may be added in a variety of forms, as salts or preferably as colloidal suspensions or sols. For example, alumina sols made from aluminum chloride, acetate, or nitrate are preferred sources of the alumina component. Readily available silica sols are preferred sources of the silica component. In each case, however, care must be taken to avoid contamination of these binder sols by elements that are harmful to the active Fischer-Tropsch metals. For example, alkali and alkaline earth cations and sulfur-containing anions such as sulfate are potent poisons of cobalt under Fischer-Tropsch conditions, and hence must be minimized in preparing supports for cobalt catalysts.

Titania supports are typically calcined after forming in order to anneal the binders, and, optionally, to convert the anatase phase of titania to the rutile phase. This calcination is typically performed in air at temperatures ranging from 500 to 1000°C.

When preparing Fischer-Tropsch catalysts from this support, metals catalytically active for the Fischer-Tropsch synthesis are composited with the support. Preferred metals are those from Group VIII of the Periodic Chart of the Elements, particularly iron, cobalt and ruthenium, with cobalt and ruthenium being preferred and cobalt being most preferred. Promoters may also be employed such as zirconium, titanium, rhenium, hafnium, cerium, thorium and uranium, and others well known to

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those skilled in the art. The metal or metals are present in amounts that are catalytically active for Fischer-Tropsch synthesis and will vary with the metal being selected. For example, ruthenium is much more active in this environment than cobalt and, as a consequence is used in amounts ranging from about 0.5-3.0 wt% while cobalt will preferably be used in amounts of about 2-40 wt%, more preferably 5-30 wt%, still more preferably 10-25 wt%.

When promoters are employed, they are used in quantities less than the active catalytic metal, e.g., in weight ratios of about 1/20 to 1/10 based on the active metal. (This invention also contemplates the use of ruthenium as a promoter in conjunction with cobalt as the primary active catalytic metal.) The most preferred catalysts are those containing cobalt and rhenium, cobalt and ruthenium, and cobalt and thoria, particularly cobalt and rhenium.

The catalyst can be prepared by a variety of techniques well known to those skilled in the art, including impregnation (either co-impregnation with promoters or serial impregnation -- either dry or by the incipient wetness techniques). Since a preferred catalyst for fixed bed Fischer-Tropsch processes is one wherein the catalytic metals are present in the outer portion of the catalyst particle, i.e., in a layer no more than 250 microns deep, preferably no more than 200 microns deep, a preferred method of preparing the catalyst is the spray method which is described in U.S. Patent 5,140,050, incorporated herein by reference or in EP 0,266,898, incorporated herein by reference. For slurry Fischer-Tropsch processes, catalysts are preferably made by incipient wetness impregnation of spray-dried supports.

Measurement of the strength of titania containing particles is not an easy task because finely divided attrition products have a tendency to adhere to the original particles and are not detectable by conventional methods, e.g., the well known Davison Attrition Test, Microtrac or Malvern light diffraction instruments. Consequently, a new accelerated attrition test was developed, wherein the results are reproducible, and based

on SEM photo micrographs of attrited products is believed to operate in an erosive or grinding environment similar to the environment in large slurry, bubble column units.

The test consists of: treatment of a small sample, e.g., about 0.5 to about 3 grams, of catalyst or support suspended in acetone, and subjected to an ultrasonic bath for a prescribed period of time, and followed by filtration to quantify the amount of fines, i.e., particles less than 25 microns mean diameter, produced. Thus, 2.5 grams of sample, sieved to +45 micron size, and suspended in 12 ml acetone is charged to a 0.5 oz. bottle. After sonicating in a Branson Model 2200 ultrasonic bath, typically for thirty minutes or more, the mixture is filtered through a 500 mesh screen (25 micron openings) onto a 0.02 micron filter membrane (Whatman Anodisc). The solids are then washed with acetone, dried and weighed to determine the weight percent (wt%) <25 micron yield.

The Fischer-Tropsch synthesis is a well known process and the reaction conditions have been described in the available literature. For example, temperatures may range from about 175° to about 400°C, preferably about 180-250°C while pressures may range from about 1-100 bar, preferably about 15 to 40 bar. Hydrogen/CO ratios may range from 0.5/1 to about 4/1, preferably about 1.7/1 to 2.5/1, with the stoichiometric amount plus or minus about 3% being most preferred. The catalyst made from the binder of this invention is preferably used in a slurry, e.g., a slurry bubble column, reactor where gas hourly space velocities may range from about 4000 to 20000. A preferred slurry bubble column operation is described in U.S. Patent #5,348,982 incorporated herein by reference.

EXAMPLES

Twelve titania supports were prepared by spray-drying mixtures of various binders with Degussa P-25 TiO₂. Dried supports were calcined between 700°C and 1000°C in rotary calciners. The amount and source of the alumina binder and of the

silica binder, the wt% solids in the spray-drier feed, and the final calcination temperature, used for each of these twelve supports are summarized in Table 1. The alumina chlorhydrol sol was made by GRACE Davison, designated as CX-100, and contained about 23.5 wt% Al₂O₃. Analytical inspections are also shown in Table 1, including the data from 30 minute sonic attrition tests. The rutile content refers to the weight percent of the rutile phase in the titania, with the balance being the anatase phase, determined by X-ray diffraction (ASTM D 3720-78). SA refers to the BET surface area and PV refers to the pore volume of pores less than about 5000 angstroms in diameter, measured by mercury porosimetry (using a mercury contact angle of 125 degrees).

Examples 1 & 2 represent the base case with an alumina binder and possess reasonably good strength here but, as will be shown later, lose strength upon steaming. Examples 3-5 with silica sols from different commercial suppliers produced very weak particles with a high portion of fines present after calcining, indicating that the use of silica binder alone is not a viable option. Example 6 illustrates this invention with the incorporation of a binder consisting of a mixture of alumina and silica sols in 9-to-1 weight ratio. Already this support shows the lowest sonic attrition value, and hence best strength, among these unsteamed examples.

Examples 7-12 in Table 1 were calcined at 700°C and hence contain lower rutile content in the titania, higher surface area, and higher pore volume compared to Examples 1-6. The higher pore volume is an especially important feature because more active metal can be deposited per incipient wetness impregnation as pore volume increases. Example 7, using the same mixture of alumina and silica binders as Example 6, illustrates that the significant improvement in pore volume gained at the lower calcination temperature can be achieved with only minor loss in particle strength. Comparing Example 8 with 7 indicates that changing the proportions of alumina to silica from 9:1 to 2:1, in a constant 6 wt% total binder composition, results in a good pore volume. However, higher total binder concentration, 12 wt% versus 6 wt%, gives lower pore volume, as shown by Examples 8, 9, and 12. Examples 11 and 12 illustrate that

very similar results are obtained by using aluminum nitrate instead of alumina chlorhydrol sol as the source of the alumina binder.

Table 2 summarizes the inferior results obtained when silica is added to alumina-bound supports *after* spray-drying. Silica was added by impregnation of tetraethyl silicate from methanol solution, followed by drying and calcining at 800°C for 3 hours in a lab oven. As indicated in the table, much of the silicate added was lost by volatilization during the drying or calcining, which poses a serious problem for this method. More importantly, the silica that remained on the support did nothing to improve particle strength. In Example 13, addition of silica to the support of Example 1 failed to noticeably improve strength. Example 14 used a support with the same composition as Example 1, but was a much weaker starting material as a result of being recalcined at 850°C in a static oven. Example 15 used a very weak support prepared by spray-drying an experimental, precipitated form of titania instead of Degussa P-25. In both Examples 14 & 15, the supports became even weaker after the silica addition, reflecting perhaps further weakening of the alumina binder by steaming in the calcination step. From these examples it is clear that for silica to function together with alumina as an improved binder, the two must be present in the initial spray-drying step.

To further define credits for the new SiO₂-Al₂O₃ binder, two Co-Re catalysts were prepared: a base case with a titania support containing 6% Al₂O₃ binder, and an example of this invention comprising a titania support with 6% of 9:1 Al₂O₃-SiO₂ binder. The spray-dried supports were made similarly as in Examples 1 and 6 but on a larger scale. Each catalyst was then made by incipient wetness impregnation of an aqueous solution of cobalt nitrate and perrenic acid, followed by air calcination at about 400°C in a rotary calciner. Double impregnation/calcinations were applied to achieve the final metal loadings. The catalysts were tested with 2/1, H₂/CO synthesis gas in a small fixed bed reactor, after reduction at 375°C. Both catalysts were highly active and selective for the formation of hydrocarbons, as shown in Table 3.

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The final, most critical strength test comes when subjecting the catalyst to high steam partial pressure at moderate temperature, such as that generated in hydrocarbon synthesis. To simulate this environment, portions of each catalyst were charged to a fluidized bed reactor and treated with pure steam at about 250°C, 6.8 atm pressure, for a period of six days. Fresh and steamed versions were then evaluated for attrition resistance using the sonic attrition test at a variety of sonication times. Results are summarized as Examples 16-19 in Table 4 and plotted in Figure 1. Referring to Figure 1, it is clear that the amount of fines produced in the attrition test increases with sonication times for all materials. Steaming weakens the alumina binder case, as shown by line A of data. The silica-alumina binder of this invention (line C) weakens only very slightly upon steaming, still maintaining a credit over even the unsteamed alumina case (line B). A remarkable improvement in initial strength and steam stability has been achieved with the improved binder.

Table 1: Spray-Dried Titania Supports

Example	Wt% Al ₂ O ₃	Al ₂ O ₃ Source	Wt% SiO ₂	SiO ₂ Source	Wt% Solids in Spray-Dryer Feed	Calcination Temp, C	Rutile	SA	PV	Sonic Attrition
1	6	Chlorhydrol sol	0	none	36	800	94	17	0.36	3.9
2	6	Chlorhydrol sol	0	none	36	800	95	14	0.34	3.6
3	0	none	6	Nyacol 2034 D.I.	23		83	20	0.28	--
4	0	none	6	Nalco 2327	20		87	17	0.30	--
5	0	none	6	Nalco 1034 A	20		92	17	0.30	10.5
6	5.4	Chlorhydrol sol	0.6	Nyacol 2034 D.I.	36	1000	93	17	0.33	1.1
7	5.4	Chlorhydrol sol	0.6	Nyacol 2034 D.I.	36	700	16	44	0.54	3.2
8	4	Chlorhydrol sol	2	Nyacol 2034 D.I.	36	700	14	51	0.50	--
9	10.8	Chlorhydrol sol	1.2	Nyacol 2034 D.I.		700	14	52	0.37	--
10	8	Chlorhydrol sol	4	Nyacol 2034 D.I.		700	15	55	0.38	--
11	5.4	Aluminum nitrate	0.6	Nyacol 2034 D.I.		700	14	50	0.47	--
12	10.8	Aluminum nitrate	1.2	Nyacol 2034 D.I.		700	13	55	0.35	--

**TABLE 2: ADDITION OF SILICA TO SUPPORTS WITH ALUMINA
BINDERS**

Example	Wt% SiO₂ Added	Wt% SiO₂ Found	Sonic Attrition Before SiO₂ Added	Sonic Attrition After SiO₂ Added
13	0.6	0.13	3.9	3.2
14	0.6	0.13	14.7	30.4
15	0.6	0.21	37.6	46.1

**TABLE 3: HCS TESTS OF CATALYSTS WITH ALUMINA AND SILICA-
ALUMINA BOUND SUPPORTS**

200°C, 280 psig, 64%H₂-32%CO-4%He

Binder	Al₂O₃	9:1 Al₂O₃-SiO₂
Wt% Co	12.0	10.6
Wt% Re	1.0	0.9
Bulk Density, g/cc	1.33	1.47
GHSV	3000	3000
% CO conversion	66	77
Mol% CH₄	6.6	5.6

TABLE 4: EFFECT OF STEAMING ON CATALYST ATTRITION

Example	Binder	Treat	Sonic Attrition Test	
			Minutes	Wt% 25-
16	Al ₂ O ₃	Fresh	0	0.2
			10	0.4
			20	0.7
			20	0.9
			30	3.6
			40	3.8
			60	8.5
			90	7.0
			90	6.3
17	Al ₂ O ₃	Steamed 6 days	0	0.7
			10	5.0
			20	5.4
			20	6.2
			30	7.4
			30	12.0
			40	11.6
			60	18.5
			90	22.1
			120	28.4
18	SiO ₂ -Al ₂ O ₃	Fresh	30	0.4
			30	0.5
			60	1.3
			120	1.1
			120	1.9
19	SiO ₂ -Al ₂ O ₃	Steamed 6 days	30	1.1
			30	1.4
			30	2.3
			60	2.7
			60	4.5
			120	6.6
			120	8.8

CLAIMS:

1. A catalyst support composition comprising primarily titania with which there is incorporated a binder comprised of silica and alumina.
2. The support of claim 1 wherein the amount of binder is less than about 30 wt% of the support and silica is less than about 50 wt% of the binder.
3. The support of claim 2 wherein the support is at least about 80 wt% titania.
4. The support of claim 2 wherein the binder is about 3-20 wt% of the support and silica is about 3-50 wt% of the binder.
5. The support composition of claim 1 wherein a metal active for promoting a Fischer-Tropsch reaction is dispersed on the support.
6. The composition of claim 5 wherein the metal is a Group VIII metal.
7. The composition of claim 6 wherein the metal is selected from the group consisting of cobalt, ruthenium, and mixtures thereof.
8. The composition of claim 6 wherein the metal is promoted with a metal selected from the group consisting of rhenium, hafnium, zirconium, cerium, thorium, and manium.
9. A Fischer-Tropsch process which comprises reacting hydrogen and carbon monoxide over the composition of claim 5 and recovering C₅+ hydrocarbons.

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10. The process of claim 9 conducted over the composition of claim 7.
11. The process of claim 9 conducted in a slurry.



FIG. 1

INTERNATIONAL SEARCH REPORT

International Application No

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A. CLASSIFICATION OF SUBJECT MATTER
IPC 6 B01J21/08 C10G2/00 C07C1/04

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IPC 6 B01J C10G C07C

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Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	GB 2 070 959 A (MATSUSHITA ELECTRIC IND CO LTD) 16 September 1981 see claims 1,2,7,10 see page 5, line 4 - line 12 ---	1,5-7
X	US 5 215 953 A (ENGLER BERND ET AL) 1 June 1993 see claim 1 ---	1
A		1,3
A	US 5 348 982 A (HERBOLZHEIMER ERIC ET AL) 20 September 1994 see claims 1,4-6 see column 5, line 54 - column 6, line 10 ---	1,5-11
A	EP 0 266 898 A (EXXON RESEARCH ENGINEERING CO) 11 May 1988 cited in the application see claims 1-4,6,10 ---	1,5-11
	-/--	

☒ Further documents are listed in the continuation of box C.

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INTERNATIONAL SEARCH REPORT

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C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category ²	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 4 992 406 A (MAULDIN CHARLES H ET AL) 12 February 1991 see claims 1,3,5-8,10,12-14,16 see table 1 ---	1,3,5-11
A	US 5 140 050 A (MAULDIN CHARLES H ET AL) 18 August 1992 cited in the application see claims 1-6 see table 1 -----	1,3,5-11

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 99/02540

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
GB 2070959 A	16-09-1981	JP 1398865 C	07-09-1987
		JP 56126447 A	03-10-1981
		JP 62001303 B	12-01-1987
		JP 1350934 C	28-11-1986
		JP 56126446 A	03-10-1981
		JP 61015733 B	25-04-1986
		JP 1320700 C	11-06-1986
		JP 57027139 A	13-02-1982
		JP 60043169 B	26-09-1985
		CA 1163615 A	13-03-1984
		US 4350613 A	21-09-1982
US 5215953 A	01-06-1993	DE 4018324 C	06-02-1992
		CA 2044067 A	09-12-1991
		CN 1057212 A,B	25-12-1991
		DE 59102409 D	08-09-1994
		EP 0460388 A	11-12-1991
		JP 4227056 A	17-08-1992
		RU 2047350 C	10-11-1995
US 5348982 A	20-09-1994	AU 632413 B	24-12-1992
		AU 7401591 A	10-10-1991
		CA 2038774 A	05-10-1991
		DE 69104400 D	10-11-1994
		DE 69104400 T	04-05-1995
		EP 0450860 A	09-10-1991
		NO 180001 B	21-10-1996
EP 0266898 A	11-05-1988	AU 623756 B	21-05-1992
		AU 5220890 A	03-10-1991
		AU 589806 B	19-10-1989
		AU 7930687 A	14-04-1988
		CA 1312066 A	29-12-1992
		CA 2012682 A	21-09-1991
		DE 3750154 D	04-08-1994
		DE 3750154 T	20-10-1994
		EP 0453674 A	30-10-1991
		JP 2572079 B	16-01-1997
		JP 63147545 A	20-06-1988
		NO 178611 B	22-01-1996
		US 4962078 A	09-10-1990
		US 4977126 A	11-12-1990
		US 5545674 A	13-08-1996
		US 5128377 A	07-07-1992
US 4992406 A	12-02-1991	AU 621955 B	26-03-1992
		AU 4532989 A	31-05-1990
		CA 1337409 A	24-10-1995
		EP 0370757 A	30-05-1990
		JP 2191549 A	27-07-1990
		JP 2849134 B	20-01-1999
		NO 180572 B	03-02-1997
US 5140050 A	18-08-1992	US 5140050 A	18-08-1992
		US 4992406 A	12-02-1991
		AU 621955 B	26-03-1992
		AU 4532989 A	31-05-1990
		CA 1337409 A	24-10-1995

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 99/02540

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US 5140050 A		EP 0370757 A	30-05-1990
		JP 2191549 A	27-07-1990
		JP 2849134 B	20-01-1999
		NO 180572 B	03-02-1997
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